

[illegible]

- wherein SP represents any sequence; SA represents a splice acceptor; SD represents a splice

donor; IRES represents an internal ribosomal entry site; M represents a marker gene; *puro* represents ; pA represents a poly(A) sequence; and PV represents a plasmid vector.

8. The trap vector of claim 7, wherein the plasmid vector is any one selected from the group consisting of pBR, pUC, pSP and pGEM.

9. A vector generated from recombination between the trap vector of claim 1 and the trap vector of claim 4.

10. The vector of claim 9, wherein said vector does not undergo recombination with other *loxP*.

11. A method of gene trapping, comprising introducing the trap vector of any one of claims 1 to 8 into embryonic stem cells.

12. Embryonic stem cells into which the trap vector of any one of claims 1 to 8 is introduced.

13. A transgenic animal into which the trap vector of any one of claims 1 to 8 is introduced.

14. The transgenic animal of claim 13, wherein said animal is selected from the group consisting of mouse, rat, rabbit, guinea pig, pig, sheep and goat.

15. A method for producing a transgenic animal, comprising introducing the embryonic stem cells of claim 12 into an animal.

16. A knockout animal into which the trap vector of any one of claims 1 to 8 is introduced.

17. The knockout animal of claim 16, wherein said animal is selected from the group consisting of mouse, rat, rabbit, guinea pig, pig, sheep and goat.

18. A method for producing a knockout animal, comprising introducing the embryonic stem cells of claim 12 into an animal.